

quantities of the  $D_2O$  may be increased. A typical example utilizing the method according to the present invention is as follows:

Within the range of between about  $20^\circ$  and  $200^\circ$  C. and at pressures of about one atmosphere to about 200 atmospheres, up to  $33 \frac{1}{2}$  atom percent  $\approx 1$  percent by weight of hydrogen are incorporated as TiNiH in the TiNi alloy. In contrast thereto under the same condition no dissolution of deuterium takes place in the TiNi-lattice, as can be readily proved. Hydrogen consists to about 99.9855% of the isotope  $^1H$  and to about 0.0145% of the isotope  $^2D$ . Consequently, 0.145 mol of deuterium correspond to 1,000 mol of hydrogen each. If one causes the content of a 100 liter pressure container with a filling pressure of 224 atmospheres (1,000 mol hydrogen) to react with the TiNi alloy at temperatures at which the dissociation pressure of the hydrogen amounts to at most 2 atmospheres, then 1,000 mol hydrogen are dissolved in 200 kg TiNi (50 l volume) within the temperature range between about  $20^\circ$  C. and about  $250^\circ$  C. If now the pressure container is now disconnected at an interior pressure of one atmosphere, then only about 5 mol hydrogen remain in the container whereas 0.145 mol deuterium remain in the same since this was not absorbed in the TiNi-lattice. If this residual gas is compressed to a volume of 500  $cm^3$  (approximately 220 atmospheres) and is brought into contact under the conditions indicated hereinabove with 1.5 kg TiNi, then the residual gas resulting therefrom consists at about 5 atmospheres of 0.02 mol hydrogen and 0.145 mol deuterium.

The speed of separation depends only on the cooling speed of the TiNi alloy (reaction enthalpy smaller than 10 k cal/mol  $H_2$ ), it amounts with the aforementioned dimensions and with a sufficient water cooling to about 15 to 30 minutes.

Consequently, in only two stages the ratio H:D of 7000:1 can be changed into 1:7, i.e., by about a factor of 50,000. A greater enrichment or concentration can be realized either by way of further hydride stages or with the aid of other known methods (Clusius separating column). In that connection the thermal separating effect can possibly be further assisted by the absorption of the hydrogen in the TiNi alloy. Consequently, depending on the desired efficiency, any required quantity of the TiNi alloy can be brought into the respectively desired temperature zone in the Clusius column.

The powder particle sizes is not uniform and may be selected between about  $10\mu$  and about 1 mm.

The mixtures may be between 0-100% and 100-0% of  $Ti_2Ni$  and TiNi.

The dehydrogenation takes place at pressures of about 0.1 atmosphere beginning with temperatures of about  $100^\circ$  C. and with greater velocity beginning with temperatures of about  $200^\circ - 250^\circ$  C.

Up to about 10% by weight of zirconium may be used instead of titanium for reducing the binding enthalpy of the hydrogen.

Up to about 25% by weight of copper may be used instead of nickel to produce a better mechanical stability of the individual granules. Nonetheless, the porosity and form of the sintered bodies remain preserved.

In the drawing the single FIGURE schematically illustrates an installation which may be used for carrying out the present invention.

Reference numeral 10 thereby generally designates a conventional pressure vessel or container having a piston or plunger 11 which is adapted to be actuated by

conventional means. Hydrogen is fed into the vessel 10 at 224 atmospheres through inlet line 12 by way of control valve 13. The pressure container 10 is connected, on the one hand, by way of a line 14 containing a control valve 15 with a second container or vessel 16 which contains TiNi powder, for example, 200 kg (50 l) of the alloy. The vessel 16 is appropriately heated or cooled by the use of conventional means, such as an appropriate conventional heating and cooling system generally designated by reference numeral 17 and therefore only schematically indicated. The discharge or removal from the container 16 takes place by way of a line 18 including a control valve 19.

The pressure container 10 is further connected with a small vessel or container 20 by way of a line 21 containing a control valve 22. The container 20 is again adapted to be heated and cooled by conventional means only schematically indicated and designated generally by reference numeral 23. For example, about 1.5 kg of the TiNi powder is provided in the container 20. The outlet from the container 20 takes place by way of line 24 including the control valve 25.

A further line 26 including a control valve 27 is provided to permit the removal of  $D_2$  and  $H_2$  for further enrichment of concentration.

In the position 1 of the piston 11, valves 15, 19, 22 and 27 are closed while valve 13 is opened. The pressure in the 100 liter vessel or container 10 is 224 atmospheres. The valve 13 is then closed and valve 15 is opened. The hydrogen then reacts with the TiNi alloy in the container 16. The pressure in the container 10 is now permitted to become about 1 atmosphere and valve 15 is closed.

The piston 11 is now moved to its position 2 indicated in dash lines whereupon the valve 19 is opened, the heating system 17 is controlled to increase the temperature by about  $100^\circ$  C. thereby causing a discharge of hydrogen out of the TiNi container 16.

The pressure underneath the piston 11 is about 200 atmospheres. Valve 22 is now opened and the hydrogen reacts with the TiNi powder in the container 20. The pressure under the piston 11 is thereby permitted to become about 5 atmospheres. Valve 22 is now closed and valve 25 is opened while the heating system 23 is controlled to provide an increase of about  $100^\circ$  C. causing a discharge of  $H_2$ . Thereupon the heating system 17 is controlled to reduce the temperature by about  $100^\circ$  C., valve 27 is opened until the pressure under the piston reaches about  $10^{-2}$  Torr. Valve 27 and valve 25 are then closed and the heating system 23 is controlled to reduce the heat by about  $100^\circ$  C.

While I have shown and described only one embodiment in accordance with the present invention, it is understood that the same is not limited thereto but is susceptible of numerous changes and modifications as known to those skilled in the art and I, therefore, do not wish to be limited to the details shown and described herein but intend to cover all such changes and modifications as are encompassed by the scope of the appended claims.

What I claim is:

1. A method for separating deuterium and hydrogen, comprising the steps of bringing into contact a gaseous mixture of deuterium and hydrogen with a metal alloy material consisting essentially of TiNi,  $Ti_2Ni$ , or a mixture thereof, and thereafter separating a non-absorbed residual gas enriched in deuterium from the metal alloy material after a period of time sufficient for the absorp-